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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/523,343	08/26/2005	Wang Min	21108.0021U2	5753
	7590 08/09/2007 OSENBERG, P.C.		EXAMINER	
SUITE 1000 999 PEACHTREE STREET			PAK, YONG D	
ATLANTA, G			ART UNIT PAPER NUMBER	
,			1652	
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			08/09/2007	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)	-
· ·	10/523,343	MIN ET AL.	
Office Action Summary	Examiner	Art Unit	
	Yong D. Pak	1652	
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet w	vith the correspondence addre	ss
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA  - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period w  - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUN 36(a). In no event, however, may a rill apply and will expire SIX (6) MO cause the application to become A	CATION. reply be timely filed  NTHS from the mailing date of this comminion (35 U.S.C. § 133).	
Status			
1)⊠ Responsive to communication(s) filed on <u>09 Fe</u> 2a)□ This action is <b>FINAL</b> . 2b)□ This     3)□ Since this application is in condition for allowar closed in accordance with the practice under E	action is non-final. nce except for formal ma	•	erits is
Disposition of Claims			
4) ⊠ Claim(s) <u>1-64</u> is/are pending in the application. 4a) Of the above claim(s) is/are withdray 5) □ Claim(s) is/are allowed. 6) □ Claim(s) is/are rejected. 7) □ Claim(s) is/are objected to. 8) ⊠ Claim(s) <u>1-64</u> are subject to restriction and/or expressions.			
Application Papers			
9) The specification is objected to by the Examiner 10) The drawing(s) filed on is/are: a) access Applicant may not request that any objection to the of Replacement drawing sheet(s) including the correction of the original transfer and the correction is objected to by the Examiner	epted or b)  objected to drawing(s) be held in abeya ion is required if the drawing	nce. See 37 CFR 1.85(a). g(s) is objected to. See 37 CFR 1	
Priority under 35 U.S.C. § 119		•	
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of:  1. Certified copies of the priority documents 2. Certified copies of the priority documents 3. Copies of the certified copies of the priority application from the International Bureau * See the attached detailed Office action for a list of	s have been received. s have been received in a ity documents have been (PCT Rule 17.2(a)).	Application No n received in this National Sta	ige
	·	•	
Attachment(c)			
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	Paper No	Summary (PTO-413) (s)/Mail Date Informal Patent Application	

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## **DETAILED ACTION**

This application is a 371 of PCT/US03/22847.

Claims 1-64 are pending.

## Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-10, drawn to a mutant thioredoxins which is resistant to oxidizing effects of cytokines, reactive oxygen species or S-nitrosylation of a SH group by nitrous oxide.

Group II, claim(s) 11-14 and 17(in part), drawn to a method of decreasing inflammation by using the thioredoxins of Group I.

Group III, claim(s) 15 and 17(in part), drawn to a method of decreasing apoptosis by using the thioredoxins of Group I.

Group IV, claim(s) 16 and 17(in part), drawn to a method of decreasing insulin resistance by using the thioredoxins of Group I.

Group V, claim(s) 18-22, drawn to a method of treating atherosclerosis using the thioredoxins of Group I.

Group VI, claim(s) 23-28, drawn to a method of treating a subject with diabetes using the thioredoxins of Group I.

Group VII, claim(s) 29-34, drawn to a method of treating a subject with an apoptotic disease using the thioredoxins of Group I.

Group VIII, claim(s) 35-40, drawn to a method of treating a subject with cardiac dysfunction using the thioredoxins of Group I.

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Group IX, claim(s) 41-48, drawn to a method of treating a subject with an angiogenesis-dependent disease using the thioredoxins of Group I.

Group X, claim(s) 49-53, drawn to a method of diagnosing an angiogenesis dependent disease using the thioredoxins of Group I.

Group XI, claim(s) 54-59, drawn to a method of screening a subject for a genetic risk of an angiogenesis dependent disease using the thioredoxins of Group I.

Group XII, claim(s) 60-64, drawn to a method of screening a subject for a genetic risk for an apoptotic disease using the thioredoxins of Group I.

The inventions listed as Groups I-XII do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The technical feature linking Groups I-XII appears to be that they all relate to a mutant thioredoxins which is resistant to oxidizing effects of cytokines, reactive oxygen species or S-nitrosylation of a SH group by nitrous oxide.

However, Bishopric et al. (form PTO-1449) discloses a thioredoxin comprising a substitutuion at a cysteine residue at position 32 and 35. Examiner takes the position that the mutant thioredoxins of Bishopric et al. inherently possesses the same material structure and functional characteristics as the enzyme of the instant invention since (1) both enzymes have the same amino acid substitutions (at residues 32 and 35), (2) both enzymes function as thioredoxins with decreased oxidizing effects, and (3) the Office does not have facilities for examining and comparing applicant's enzyme with the enzyme of the prior art, the burden is on the applicant to show a novel or unobvious

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difference between the claimed product and the product of the prior art (i.e., that the mutant thioredoxins of the prior art does not possess the same material structure and functional characteristics of the claimed thioredoxin). See *In re Best*, 562 F.2d 1252, 195 USPQ 430 (CCPA 1977) and *In re Figzgerald* et al., 205 USPQ 594.

Therefore, the technical feature linking the inventions of Groups I-XII does not constitute a special technical feature as defined by PCT Rule 13.2, as it does not define a contribution over the prior art.

The special technical feature of Group I is a mutant thioredoxins which is resistant to oxidizing effects of cytokines, reactive oxygen species or S-nitrosylation of a SH group by nitrous oxide.

The special technical feature of Group II is a method of decreasing inflammation by using a thioredoxin.

The special technical feature of Group III is a method of decreasing apoptosis by using a thioredoxin.

The special technical feature of Group IV is a method of decreasing insulin resistance by using a thioredoxin.

 The special technical feature of Group V is a method of treating atherosclerosis using a thioredoxin.

The special technical feature of Group VI is a method of treating a subject with diabetes using a thioredoxin.

The special technical feature of Group VII is a method of treating a subject with an apoptotic disease using a thioredoxin.

The special technical feature of Group VIII is a method of treating a subject with cardiac dysfunction using a thioredoxin.

The special technical feature of Group IX is a method of treating a subject with an angiogenesis-dependent disease using a thioredoxin.

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The special technical feature of Group X is a method of diagnosing an angiogenesis dependent disease using a thioredoxin.

The special technical feature of Group XI is a method of screening a subject for a genetic risk of an angiogenesis dependent disease using a thioredoxin.

The special technical feature of Group XII is a method of screening a subject for a genetic risk for an apoptotic disease using a thioredoxin.

Further, the products of Groups II-XII do not share a technical feature for the following reasons. The methods of Groups II-XII do not share special technical features because the methods have different steps and effects.

Accordingly, Groups I-XII are not so linked by the same or a corresponding special technical feature as to form a single general inventive concept.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Yong Pak whose telephone number is 571-272-0935. The examiner can normally be reached 6:30 A.M. to 5:00 P.M. Monday through Thursday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Ponnathapu Achutamurthy can be reached on 571-272-0928. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 571-272-1600.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll free).

Yong D. Pak

Patent Examiner 1652